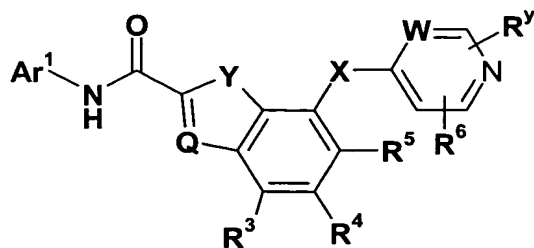


**What is Claimed is:**

1. A compound of the formula (I)



(I);

wherein:

$Ar^1$  is an aromatic carbocycle substituted with one  $R^1$ , and wherein  $Ar^1$  is independently substituted with two  $R^2$  groups and wherein one  $R^1$  and one  $R^2$  on adjacent ring atoms optionally form a 5- or 6-membered carbocyclic or heterocyclic ring;

$R^1$  is halogen,  $NO_2$ ,  $NH_2$ ,  $J-N(R^a)-(CH_2)_m$ ,  $N(J)_2-(CH_2)_m$ ,  $NH_2C(O)-$ ,  $J-N(R^a)-C(O)-$ ,  $J-S(O)_m-N(R^a)-$ ,  $J-N(R^a)-S(O)_m$ - or heterocycle- $(CH_2)_m$ - wherein the heterocyclic group is optionally substituted by  $C_{1-5}$  alkyl;

$Q$  is a  $N$  or  $CR^p$ ;

$Y$  is  $>CR^pR^v$ ,  $-CR^p=C(R^v)-$ ,  $-O-$ ,  $-N(R^x)-$  or  $>S(O)_m$ ;

wherein  $R^a$ ,  $R^p$ ,  $R^v$ ,  $R^x$  and  $R^y$  are each independently hydrogen or  $C_{1-5}$  alkyl;

$X$  is  $-CH_2-$ ,  $-N(R^a)-$ ,  $-O-$  or  $-S-$ ;

$W$  is  $N$  or  $CH$ ;

each  $m$  is independently 0, 1 or 2;

$J$  is chosen from  $C_{1-10}$  alkyl and carbocycle each optionally substituted by  $R^b$ ;

**R<sup>2</sup>** is chosen from C1-6 alkyl, C3-7 cycloalkyl optionally substituted by C1-5 alkyl, C1-4 acyl, aroyl, C1-4 alkoxy, each being optionally partially or fully halogenated, halogen, C1-6 alkoxycarbonyl, carbocyclesulfonyl and -SO<sub>2</sub>-CF<sub>3</sub>;

5

each **R<sup>3</sup>**, **R<sup>4</sup>** and **R<sup>5</sup>** are independently chosen from hydrogen, C1-6 alkyl and halogen;

**R<sup>6</sup>** is optionally attached at a position *ortho* or *meta* to the N atom of the indicated ring, and is chosen from

- 10 a bond, -O-, -O-(CH<sub>2</sub>)<sub>1-5</sub>-, >C(O), -NH-, -C(O)-NH-, -S-, C<sub>1-5</sub> alkyl branched or unbranched, C<sub>2-5</sub> alkenyl, C<sub>1-3</sub> acyl, C<sub>1-3</sub> alkyl(OH), heterocycle selected from morpholinyl, piperazinyl, piperidinyl, pyrrolidinyl and tetrahydrofuranyl, heteroaryl selected from pyridinyl, pyrimidinyl, pyrazinyl, pyridazinyl, pyrrolyl, imidazolyl, pyrazolyl, thienyl, furyl, isoxazolyl, thiazolyl, oxazolyl and isothiazolyl or aryl each
- 15 alkyl, alkenyl, acyl, heterocycle, heteroaryl and aryl are optionally substituted by one to three hydroxy, oxo, C<sub>1-3</sub> alkyl, C<sub>1-3</sub> alkoxy, C<sub>1-5</sub> alkoxycarbonyl, -NR<sub>7</sub>R<sub>8</sub> or NR<sub>7</sub>R<sub>8</sub>-C(O)-;

wherein each **R<sub>6</sub>** is further optionally covalently attached to groups chosen from:

- 20 hydrogen, -NR<sub>7</sub>R<sub>8</sub>, C<sub>1-3</sub> alkyl, C<sub>3-6</sub> cycloalkylC<sub>0-2</sub>alkyl, hydroxy, C<sub>1-3</sub> alkoxy, phenoxy, benzyloxy, arylC<sub>0-4</sub> alkyl, heteroaryl C<sub>0-4</sub> alkyl and heterocycle C<sub>0-4</sub>alkyl, each above-listed heterocycle, heteroaryl and aryl group is optionally substituted by one to three hydroxy, oxo, C<sub>1-4</sub> alkyl, C<sub>1-3</sub> alkoxy, C<sub>1-5</sub> alkoxycarbonyl, NR<sub>7</sub>R<sub>8</sub>-C(O)- or C<sub>1-4</sub> acyl;

25

each **R<sub>7</sub>** and **R<sub>8</sub>** are independently hydrogen, phenylC<sub>0-3</sub>alkyl optionally substituted by halogen, C<sub>1-3</sub> alkyl or diC<sub>1-5</sub> alkyl amino, or **R<sub>7</sub>** and **R<sub>8</sub>** are C<sub>1-2</sub> acyl, benzoyl or C<sub>1-5</sub> branched or unbranched alkyl optionally substituted by C<sub>1-4</sub> alkoxy, hydroxy or mono or diC<sub>1-3</sub> alkyl amino;

30 and

**R<sup>b</sup>** is chosen from hydrogen, C1-5 alkyl, hydroxyC1-5 alkyl, C2-5 alkenyl, C2-5 alkynyl, carbocycle, heterocycle, heteroaryl, C1-5 alkoxy, C1-5 alkylthio, amino, C1-5 alkylamino, C1-5 dialkylamino, C1-5 acyl, C1-5 alkoxycarbonyl, C1-5 acyloxy, C1-5

acylamino, each of the aforementioned are optionally partially or fully halogenated, or  $R^b$  is chosen from C1-5 alkylsulphonylamino, hydroxy, oxo, halogen, nitro and nitrile;

or the pharmaceutically acceptable salts, acids or isomers thereof.

5

2. The compound according to claim 1 and wherein:

Y is -O-, -S-, -NH-, -N(CH<sub>2</sub>CH<sub>3</sub>)- or -N(CH<sub>3</sub>)-;

10

X is -N(R<sup>a</sup>)-, or -O-;

Q is CH;

J is chosen from C1-10 alkyl, aryl or C3-7 cycloalkyl each optionally substituted by  $R^b$ ;

15

$R_2$  is independently chosen from C1-6 alkyl, C3-6 cycloalkyl optionally substituted by C1-3 alkyl, acetyl, aroyl, C1-5 alkoxy, each being optionally partially or fully halogenated, halogen, methoxycarbonyl, phenylsulfonyl and -SO<sub>2</sub>-CF<sub>3</sub>;

20 each  $R^3$ ,  $R^4$  and  $R^5$  are hydrogen;

$R^b$  is chosen from hydrogen, C1-5 alkyl, C2-5 alkenyl, C2-5 alkynyl, C3-8 cycloalkyl, C0-2 alkyl, aryl, C1-5 alkoxy, C1-5 alkylthio, amino, C1-5 alkylamino, C1-5 dialkylamino, C1-5 acyl, C1-5 alkoxycarbonyl, C1-5 acyloxy, C1-5 acylamino, C1-5 sulphonylamino, hydroxy, halogen, trifluoromethyl, nitro, nitrile

25

or  $R^b$  is chosen from; heterocycle chosen from pyrrolidinyl, pyrrolinyl, morpholinyl, thiomorpholinyl, thiomorpholinyl sulfoxide, thiomorpholinyl sulfone, dioxalanyl, piperidinyl, piperazinyl, tetrahydrofuranyl, tetrahydropyranyl, tetrahydrofuranyl, 1,3-dioxolanone, 1,3-dioxanone, 1,4-dioxanyl, piperidinonyl, tetrahydropyrimidinyl,

30

pentamethylene sulfide, pentamethylene sulfoxide, pentamethylene sulfone, tetramethylene sulfide, tetramethylene sulfoxide and tetramethylene sulfone and heteroaryl chosen from aziridinyl, thienyl, furanyl, isoxazolyl, oxazolyl, thiazolyl, thiadiazolyl, tetrazolyl, pyrazolyl, pyrrolyl, imidazolyl, pyridinyl, pyrimidinyl, pyrazinyl, pyridazinyl, pyranyl, quinoxalinyl, indolyl, benzimidazolyl, benzoxazolyl,

benzothiazolyl, benzothienyl, quinolinyl, quinazolinyl, naphthyridinyl, indazolyl, triazolyl, pyrazolo[3,4-*b*]pyrimidinyl, purinyl, pyrrolo[2,3-*b*]pyridinyl, pyrazolo[3,4-*b*]pyridinyl, tubercidinyl, oxazo[4,5-*b*]pyridinyl and imidazo[4,5-*b*]pyridinyl.

5

3. The compound according to claim 2 and wherein:

$\text{Ar}^1$  is chosen from phenyl, naphthyl, tetrahydronaphthyl, indanyl and indenyl, each  $\text{Ar}^1$  is optionally substituted with one  $\text{R}^1$ , and independently substituted with two  $\text{R}^2$  groups;

10

$\text{Y}$  is -O-, -S- or -N(CH<sub>3</sub>)-;

$\text{R}^6$  is present, and is chosen from

15

a bond, -O-, -O-(CH<sub>2</sub>)<sub>1-5</sub>-, -NH-, -C(O)-NH-, C<sub>1-5</sub> alkyl branched or unbranched, C<sub>2-5</sub> alkenyl, C<sub>1-3</sub> alkyl(OH), heterocycle selected from morpholinyl, piperazinyl, piperidinyl, pyrrolidinyl and tetrahydrofuranyl, or aryl chosen from phenyl and naphthyl, each alkyl, alkenyl, heterocycle and aryl are optionally substituted by one to three hydroxy, C<sub>1-3</sub> alkyl, C<sub>1-3</sub> alkoxy, mono or diC<sub>1-3</sub> alkyl amino, amino or C<sub>1-5</sub> alkoxycarbonyl;

20

wherein each  $\text{R}_6$  is further optionally covalently attached to groups chosen from:

hydrogen, -NR<sub>7</sub>R<sub>8</sub>, C<sub>1-3</sub> alkyl, C<sub>3-6</sub> cycloalkylC<sub>0-2</sub>alkyl, hydroxy, C<sub>1-3</sub> alkoxy, phenoxy, benzyloxy, phenylC<sub>0-4</sub> alkyl, piperazinylC<sub>0-4</sub> alkyl, piperidinyl C<sub>0-4</sub>alkyl, pyrrolidinylC<sub>0-4</sub> alkyl, morpholinylC<sub>0-4</sub> alkyl, tetrahydrofuranylC<sub>0-4</sub> alkyl, triazolyl C<sub>0-4</sub>alkyl, imidazolyl C<sub>0-4</sub>alkyl and pyridinyl C<sub>0-4</sub>alkyl, each abovelisted heterocycle, heteroaryl and phenyl group is optionally substituted by one to three hydroxy, oxo, C<sub>1-4</sub> alkyl, C<sub>1-3</sub> alkoxy, C<sub>1-5</sub> alkoxycarbonyl, -NR<sub>7</sub>R<sub>8</sub>, NR<sub>7</sub>R<sub>8</sub>-C(O)- or C<sub>1-4</sub> acyl;

25

each  $\text{R}_7$  and  $\text{R}_8$  are independently hydrogen, phenylC<sub>0-3</sub>alkyl optionally substituted by

30

halogen, C<sub>1-3</sub> alkyl or diC<sub>1-5</sub> alkyl amino, or  $\text{R}_7$  and  $\text{R}_8$  are C<sub>1-2</sub> acyl, benzoyl or C<sub>1-5</sub> branched or unbranched alkyl optionally substituted by C<sub>1-4</sub> alkoxy, hydroxy or mono or diC<sub>1-3</sub> alkyl amino.

4. The compound according to claim 3 and wherein:

**X** is -O-;

5 **Y** is -N(CH<sub>3</sub>)-;

**J** is C1-10 alkyl optionally substituted by **R<sup>b</sup>**;

**R<sub>2</sub>** is independently chosen from C1-6 alkyl, C3-6 cycloalkyl optionally substituted by  
10 C1-3 alkyl and C1-5 alkoxy, each being optionally be partially or fully halogenated;

**R<sup>6</sup>** is chosen from  
a bond, -O-, -O-(CH<sub>2</sub>)<sub>1-5</sub>-, -NH-, -C(O)-NH-, C<sub>1-5</sub> alkyl branched or unbranched, C<sub>2-5</sub>  
alkenyl, C<sub>1-3</sub> alkyl(OH), heterocycle selected from morpholinyl, piperazinyl, piperidinyl  
15 and pyrrolidinyl or phenyl, each alkyl, alkenyl, heterocycle and phenyl are optionally  
substituted by one to three hydroxy, C<sub>1-3</sub> alkyl, C<sub>1-3</sub> alkoxy, mono or diC<sub>1-3</sub> alkyl amino,  
amino or C<sub>1-5</sub> alkoxycarbonyl;

wherein each **R<sub>6</sub>** is further optionally covalently attached to groups chosen from:  
20 hydrogen, -NR<sub>7</sub>R<sub>8</sub>, C<sub>1-3</sub> alkyl, C<sub>3-6</sub> cycloalkylC<sub>0-2</sub>alkyl, benzyloxy, phenylC<sub>0-4</sub>  
alkyl, piperazinylC<sub>0-4</sub> alkyl, piperidinyl C<sub>0-4</sub>alkyl, pyrrolidinylC<sub>0-4</sub> alkyl, morpholinylC<sub>0-4</sub>  
alkyl, triazolyl C<sub>0-4</sub>alkyl, imidazolyl C<sub>0-4</sub>alkyl and pyridinyl C<sub>0-4</sub>alkyl, each above-  
listed heterocycle, heteroaryl and phenyl group is optionally substituted by one to three  
hydroxy, oxo, C<sub>1-4</sub> alkyl, C<sub>1-3</sub> alkoxy, C<sub>1-5</sub> alkoxycarbonyl, amino, NR<sub>7</sub>R<sub>8</sub>-C(O)- or C<sub>1-4</sub>  
25 acyl;

each **R<sub>7</sub>** and **R<sub>8</sub>** are independently hydrogen, phenylC<sub>0-2</sub>alkyl optionally substituted by  
halogen, C<sub>1-3</sub> alkyl or diC<sub>1-5</sub> alkyl amino, or **R<sub>7</sub>** and **R<sub>8</sub>** are C<sub>1-5</sub> branched or unbranched  
alkyl optionally substituted by C<sub>1-4</sub> alkoxy, hydroxy or mono or diC<sub>1-3</sub> alkyl amino;

30 **R<sup>b</sup>** is chosen from hydrogen, C1-5 alkyl, C3-7 cycloalkylC<sub>0-2</sub> alkyl, aryl, C1-5 alkoxy,  
amino, C1-5 alkylamino, C1-3 dialkylamino, C1-3 acyl, C1-5 alkoxycarbonyl, C1-3

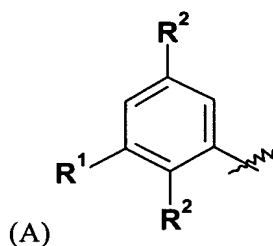
acyloxy, C1-3 acylamino, C1-3 sulphonylamino, hydroxy, halogen, trifluoromethyl, nitro, nitrile;

or  $R^b$  is chosen from pyrrolidinyl, pyrrolinyl, morpholinyl, thiomorpholinyl, thiomorpholinyl sulfoxide, thiomorpholinyl sulfone, piperidinyl, piperazinyl,

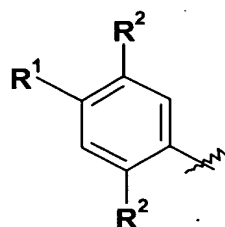
- 5 piperidinonyl, tetrahydropyrimidinonyl, aziridinyl, isoxazolyl, oxazolyl, thiazolyl, thiadiazolyl, tetrazolyl, pyrazolyl, pyrrolyl, imidazolyl, pyridinyl, pyrimidinyl, pyrazinyl and pyridazinyl.

- 10 5. The compound according to claim 4 and wherein:

$Ar^1$  is formula (A) or (B)



or (B)



- 15 wherein:

when  $Ar^1$  is formula (A) then:

$R^1$  is  $NH_2$ ,  $J-N(R^a)-(CH_2)_m-$ ,  $NH_2C(O)-$ ,  $J-N(R^a)-C(O)-$ ,  $J-S(O)_2-N(R^a)-$ ,  $J-N(R^a)-S(O)_2-$  or heterocycle- $(CH_2)_{1-2}-$  wherein the heterocycle is chosen from pyrrolidinyl, morpholinyl and piperazinyl each optionally substituted by C1-4 alkyl, and

- 20  $J$  is  $C_{1-5}$  alkyl optionally substituted by  $R^b$ ;

or

when  $Ar^1$  is formula (B) then:

$R^1$  is hydrogen or halogen;

- 25  $R_2$  is independently chosen from C1-5 alkyl, C3-6 cycloalkyl optionally substituted by C1-3 alkyl and C1-5 alkoxy, each being optionally partially or fully halogenated;

$R^6$  is chosen from

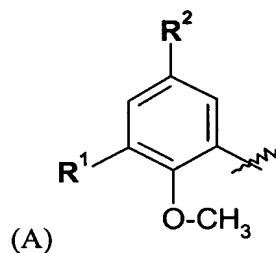
a bond, -O-, -O-(CH<sub>2</sub>)<sub>1-5</sub>-, -NH-, -C(O)-NH-, C<sub>1-5</sub> alkyl branched or unbranched, C<sub>2-5</sub> alkenyl, C<sub>1-3</sub> alkyl(OH), heterocycle selected from morpholinyl, piperazinyl, piperidinyl and pyrrolidinyl or phenyl, each alkyl, alkenyl, heterocycle and phenyl are optionally substituted by one to three hydroxy, C<sub>1-3</sub> alkyl, C<sub>1-3</sub> alkoxy, mono or diC<sub>1-3</sub> alkyl amino, amino or C<sub>1-5</sub> alkoxycarbonyl;

wherein each R<sub>6</sub> is further optionally covalently attached to groups chosen from:

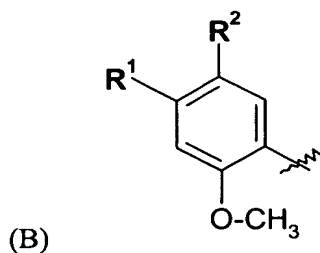
hydrogen, -NR<sub>7</sub>R<sub>8</sub>, C<sub>1-3</sub> alkyl, C<sub>3-6</sub> cycloalkylC<sub>0-2</sub>alkyl, benzyloxy, phenylC<sub>0-4</sub> alkyl, piperazinyl, piperazinylC<sub>1-2</sub> alkyl, piperidinyl, piperidinyl C<sub>1-2</sub>alkyl, pyrrolidinyl, pyrrolidinyl C<sub>1-2</sub> alkyl, morpholinyl, morpholinylC<sub>1-2</sub> alkyl, triazolyl, triazolyl C<sub>1-2</sub>alkyl, imidazolyl, imidazolyl C<sub>1-2</sub>alkyl, pyridinyl and pyridinyl C<sub>1-2</sub>alkyl, each above-listed heterocycle, heteroaryl and phenyl group is optionally substituted by one to three hydroxy, oxo, C<sub>1-4</sub> alkyl, C<sub>1-3</sub> alkoxy, C<sub>1-5</sub> alkoxycarbonyl, amino, NR<sub>7</sub>R<sub>8</sub>-C(O)- or C<sub>1-4</sub> acyl.

6. The compound according to claim 5 and wherein:

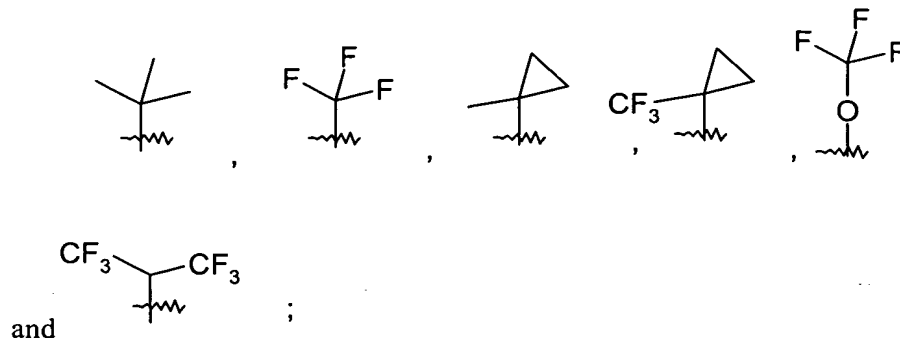
Ar<sup>1</sup> is formula (A) or (B)



or



and R<sup>2</sup> is chosen from



when  $\text{Ar}^1$  is formula (A) then:

when  $\text{R}^1$  is  $\text{J-S(O)}_2\text{-N(R}^a\text{)-}$  or  $\text{J-N(R}^a\text{)-S(O)}_2\text{-}$  then  $\text{J}$  is  $\text{C}_{1-3}$  alkyl;

and

when  $\text{R}^1$  is  $\text{NH}_2$ ,  $\text{J-N(R}^a\text{)-(CH}_2\text{)}_m\text{-}$ ,  $\text{NH}_2\text{C(O)-}$ ,  $\text{J-N(R}^a\text{)-C(O)-}$ ,

- 5 or heterocycle-( $\text{CH}_2$ )<sub>1-2</sub>- wherein the heterocycle is chosen from pyrrolidinyl, morpholinyl, piperazinyl or C1-4alkylpiperazinyl, then  $\text{J}$  is C1-3 alkyl optionally substituted by  $\text{R}^b$ .

- 10 7. The compound according to claim 6 and wherein:

$\text{R}^b$  is chosen from hydrogen, C1-5 alkyl, C3-6 cycloalkyl, C0-2 alkyl, phenyl, C1-5 alkoxy, amino, C1-5 alkylamino, C1-3 dialkylamino, C1-3 acyl, C1-5 alkoxycarbonyl, C1-3 acyloxy, C1-3 acylamino, hydroxy, halogen;

- 15 or  $\text{R}^b$  is chosen from morpholinyl, thiomorpholinyl, thiomorpholinyl sulfoxide, thiomorpholinyl sulfone, piperidinyl, piperidinonyl, pyridinyl, pyrimidinyl, pyrazinyl and pyridazinyl.

8. The compound according to claim 7 and wherein:

20

$\text{R}^b$  is chosen from amino, C1-5 alkylamino, C1-3 dialkylamino; or  $\text{R}^b$  is chosen morpholinyl, piperidinyl and pyridinyl.

9. The compound according to claim 6 and wherein:

- 25  $\text{Ar}^1$  is formula (A).

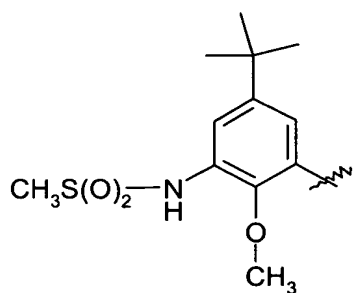
10. The compound according to claim 6 and wherein:

$\text{Ar}^1$  is formula (B).

- 30 11. The compound according to claim 6 and wherein:

$\text{Ar}^1$  is





12. A compound chosen from:

1-Methyl-7-(pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-(2-Methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-phenyl)-amide
7-[2-(2-Dimethylamino-ethylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-(2-Cyclopropylamino-pyrimidin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-[2-(4-Methoxy-benzylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(4-methyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-[2-(2-Dimethylamino-ethylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(4-methyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(2-morpholin-4-yl-ethylamino)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide

7-[2-(3-Dimethylamino-propylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-[2-(3-Dimethylamino-2,2-dimethyl-propylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-(2-Dimethylamino-pyrimidin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(6-methyl-2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(2-pyrrolidin-1-yl-ethylamino)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(piperidin-4-ylamino)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
4-{4-[2-(5-tert-Butyl-3-methanesulfonylamino-2-methoxy-phenylcarbamoyl)-1-methyl-1H-indol-7-yloxy]-pyrimidin-2-ylamino}-piperidine-1-carboxylic acid tert-butyl ester
7-{2-[(2-Dimethylamino-ethyl)-methyl-amino]-pyrimidin-4-yloxy}-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[6-methyl-2-(4-methyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-[2-(2-Dimethylamino-ethoxy)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-[2-(2-Dimethylamino-ethoxy)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(2-pyrrolidin-1-yl-ethoxy)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(2-morpholin-4-yl-ethoxy)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(1-methyl-piperidin-4-yloxy)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-[2-(2-Dimethylamino-ethoxy)-6-methyl-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide

1-Methyl-7-(2-methylcarbamoyl-pyridin-4-yloxy)-1H-indole-2-carboxylic acid [5-tert-butyl-3-(2-dimethylamino-ethylcarbamoyl)-2-methoxy-phenyl]-amide
7-[2-(2-Dimethylamino-ethylcarbamoyl)-pyridin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid [5-tert-butyl-3-(2-dimethylamino-ethylcarbamoyl)-2-methoxy-phenyl]-amide
1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid [5-tert-butyl-2-methoxy-3-(2-morpholin-4-yl-ethylcarbamoyl)-phenyl]-amide
1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-carbamoyl-2-methoxy-phenyl)-amide
1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-3-methylcarbamoyl-phenyl)-amide
1-Methyl-7-(2-vinyl-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-[2-(1,2-Dihydroxy-ethyl)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(morpholin-4-ylamino)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(2-morpholin-4-ylmethyl-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(2-morpholin-4-ylmethyl-pyridin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(4-methyl-piperazin-1-ylmethyl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-(2-Dimethylaminomethyl-pyridin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(2-methylcarbamoyl-pyridin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-(2-Benzoyloxymethyl-pyridin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide

1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-3-morpholin-4-ylmethyl-phenyl)-amide
1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid [5-tert-butyl-2-methoxy-3-(4-methyl-piperazin-1-ylmethyl)-phenyl]-amide
1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-dimethylaminomethyl-2-methoxy-phenyl)-amide
1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (3-amino-5-tert-butyl-2-methoxy-phenyl)-amide
1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-dibenzylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-3-methylsulfamoyl-phenyl)-amide
7-[2-(2-Dimethylamino-ethylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-[1,3]dioxolan-2-yl-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(4-methyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-3-methylaminomethyl-phenyl)-amide
1-Methyl-7-[2-(4-methyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-3-pyrrolidin-1-ylmethyl-phenyl)-amide
1-Methyl-7-{2-[methyl-(1-methyl-piperidin-4-yl)-amino]-pyrimidin-4-yloxy}-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-(2-Hydroxymethyl-pyridin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide and
1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid [5-tert-butyl-2-methoxy-3-(2-morpholin-4-yl-ethylamino)-phenyl]-amide
1-Methyl-7-(pyridin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(2-piperazin-1-yl-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid [3-methanesulfonylamino-2-methoxy-5-(1-methyl-cyclopropyl)-phenyl]-amide

1-Methyl-7-[2-(5-methyl-2,5-diaza-bicyclo[2.2.1]hept-2-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-[2-(2,5-Diaza-bicyclo[2.2.1]hept-2-yl)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-(2-Methoxy-pyrimidin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-[2-(4-tert-Butyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(2-morpholin-4-yl-ethyl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-{2-[2-(4-methyl-piperazin-1-yl)-ethyl]-pyrimidin-4-yloxy}-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(2-pyrrolidin-1-yl-ethyl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-[2-(2-Dimethylamino-ethyl)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(2-morpholin-4-yl-ethyl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl)-amide
1-Methyl-7-[2-(4-methyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl)-amide
7-{2-[2-(4-tert-Butyl-piperazin-1-yl)-ethyl]-pyrimidin-4-yloxy}-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-[2-(4-tert-Butyl-piperazin-1-ylmethyl)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(2-pyrrolidin-1-ylmethyl-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-(2,6-Dimethyl-pyridin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-(2-Ethyl-pyridin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide

1-Methyl-7-[2-(1,2,3,6-tetrahydro-pyridin-4-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-(2-Amino-pyrimidin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(2-pyrrolidin-1-ylmethyl-pyridin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(2-piperidin-1-ylmethyl-pyridin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(4-methyl-piperazin-1-ylmethyl)-pyridin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(pyridin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-[(2-dimethylamino-ethyl)-methyl-amino]-methyl)-2-methoxy-phenyl)-amide
7-(2-[(2-Dimethylamino-ethyl)-methyl-amino]-methyl)-pyrimidin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(4-methyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-carbamoyl-2-methoxy-phenyl)-amide
1-Methyl-7-[2-((1S,4S)-5-methyl-2,5-diaza-bicyclo[2.2.1]hept-2-yl)-pyridin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(4-methyl-[1,4]diazepan-1-yl)-pyridin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-(2-[1,4]Diazepan-1-yl-pyridin-4-yloxy)-1-methyl-1-H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(2-piperazin-1-yl-pyridin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-trideuterio-7-(2-piperazin-1-yl-pyridin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-[2-(Hexahydro-pyrrolo[1,2-a]pyrazin-2-yl)-pyridin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-trideuterio-7-[2-(4-methyl-piperazin-1-yl)-pyridin-4-yloxy]-1H-indole-2-carboxylic acid [3-methanesulfonylamino-2-methoxy-5-(1-methyl-cyclopropyl)-phenyl]-amide

7-[2-((S)-3-Dimethylamino-pyrrolidin-1-yl)-pyridin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid [3-methanesulfonylamino-2-methoxy-5-(1-methyl-cyclopropyl)-phenyl]-amide
7-[2-((S)-3-Dimethylamino-pyrrolidin-1-yl)-pyridin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid [3-methanesulfonylamino-2-methoxy-5-(1-methyl-cyclopropyl)-phenyl]-amide
1-Methyl-7-[2-(4-methyl-piperazine-1-carbonyl)-pyridin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide and
1-Methyl-7-[2-(piperazine-1-carbonyl)-pyridin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide

or the pharmaceutically acceptable salts, acids or isomers thereof.

13. A compound chosen from:

7-(Pyrimidin-4-yloxy)-benzo[b]thiophene-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-(Pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-(Pyrimidin-4-yloxy)-benzofuran-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(pyrimidin-4-ylsulfanyl)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(pyrimidin-4-ylamino)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(pyridin-3-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-(2-Benzylamino-pyrimidin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-{2-[(pyridin-2-ylmethyl)-amino]-pyrimidin-4-yloxy}-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-[2-(2-Imidazol-1-yl-ethylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(2-[1,2,3]triazol-1-yl-ethylamino)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide

7-[2-(3-Dimethylamino-propylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid [2-methoxy-5-(2,2,2-trifluoro-1-trifluoromethyl-ethyl)-phenyl]-amide
7-{2-[(2-Dimethylamino-ethyl)-methyl-amino]-pyrimidin-4-yloxy}-1-methyl-1H-indole-2-carboxylic acid (4-chloro-2-methoxy-5-trifluoromethyl-phenyl)-amide
7-[2-(4-Acetyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(4-methyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (4-chloro-2-methoxy-5-trifluoromethyl-phenyl)-amide
7-[2-(2-Dimethylamino-ethylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (2-methoxy-5-trifluoromethoxy-phenyl)-amide
7-[2-(4-Dimethylamino-piperidin-1-yl)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-[2-(3-Dimethylamino-pyrrolidin-1-yl)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(1-methyl-piperidin-4-ylamino)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-[2-(1-Acetyl-piperidin-4-ylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(2-morpholin-4-yl-ethoxy)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl)-amide
7-[2-(2-Imidazol-1-yl-ethoxy)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-[2-(2-Imidazol-1-yl-ethoxy)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (4-chloro-2-methoxy-5-trifluoromethyl-phenyl)-amide
7-[2-(2-Dimethylamino-ethylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-3-methylcarbamoyl-phenyl)-amide
7-(2-Amino-pyrimidin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-carbamoyl-2-methoxy-phenyl)-amide
7-(2-Amino-pyrimidin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid [5-tert-butyl-3-(2-dimethylamino-ethylcarbamoyl)-2-methoxy-phenyl]-amide



7-[2-(2-Dimethylamino-ethylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-dimethylaminomethyl-2-methoxy-phenyl)-amide
7-[2-(2-Dimethylamino-ethylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-3-pyrrolidin-1-ylmethyl-phenyl)-amide
7-[2-(2-Dimethylamino-ethylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-3-morpholin-4-ylmethyl-phenyl)-amide
1-Methyl-7-(2-morpholin-4-ylmethyl-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (4-chloro-2-methoxy-5-trifluoromethyl-phenyl)-amide
7-[2-(3-Dimethylamino-pyrrolidin-1-ylmethyl)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-(2-Carbamoyl-pyrimidin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-(2-morpholin-4-ylmethyl-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (2-methoxy-3-morpholin-4-ylmethyl-5-trifluoromethyl-phenyl)-amide
1-Methyl-7-[2-(4-methyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (2-methoxy-3-morpholin-4-ylmethyl-5-trifluoromethyl-phenyl)-amide
1-Methyl-7-(2-morpholin-4-ylmethyl-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl)-amide
7-(1'-tert-Butyl-1',2',3',4',5',6'-hexahydro-[2,4']bipyridinyl-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl)-amide
1-Methyl-7-(2-methylaminomethyl-pyridin-4-yloxy)-1H-indole-2-carboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl)-amide
1-Methyl-7-(2-pyrrolidin-1-ylmethyl-pyridin-4-yloxy)-1H-indole-2-carboxylic acid (2-methoxy-3-morpholin-4-ylmethyl-5-trifluoromethyl-phenyl)-amide
1-Methyl-7-[2-(2-morpholin-4-yl-ethyl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (3-dimethylaminomethyl-2-methoxy-5-trifluoromethyl-phenyl)-amide
1-Methyl-7-(2-pyrrolidin-1-ylmethyl-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (2-methoxy-3-pyrrolidin-1-ylmethyl-5-trifluoromethyl-phenyl)-amide
7-(2-Dimethylaminomethyl-pyrimidin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid [2-methoxy-3-(4-methyl-piperazin-1-ylmethyl)-5-trifluoromethyl-phenyl]-amide

7-(2-Dimethylaminomethyl-pyridin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl)-amide
7-(2-Dimethylaminomethyl-pyrimidin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl)-amide
1-Methyl-7-[2-(4-methyl-piperazin-1-ylmethyl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl)-amide
7-(2-Dimethylaminomethyl-pyrimidin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl)-amide
1-Methyl-7-[2-(2-morpholin-4-yl-ethyl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid [3-methanesulfonylamino-2-methoxy-5-(1-methyl-cyclopropyl)-phenyl]-amide
1-Methyl-7-[2-(1-methyl-piperidin-4-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
7-[2-(1-Cyclopropyl-piperidin-4-yl)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
1-Methyl-7-[2-(4-methyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (3-dimethylaminomethyl-2-methoxy-5-trifluoromethyl-phenyl)-amide and
1-Methyl-7-[2-(1-methyl-pyrrolidin-3-ylamino)-pyridin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide

or the pharmaceutically acceptable salts, acids or isomers thereof.

- 5 14. A pharmaceutical composition containing a pharmaceutically effective amount of a compound according to claim 1 and one or more pharmaceutically acceptable carriers and/or adjuvants.

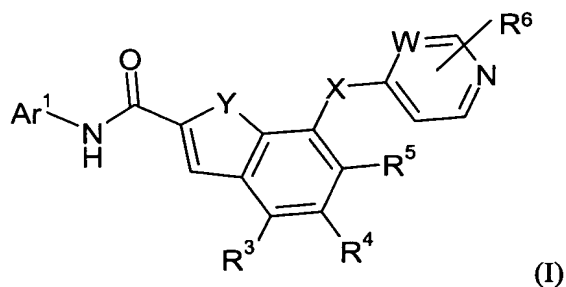
- 10 15. A method of treating an oncological disease comprising administering to a patient a pharmaceutically effective amount of a compound according to claim 1.

16. A method of treating a disease or condition chosen from osteoarthritis, atherosclerosis, contact dermatitis, bone resorption diseases, reperfusion injury, asthma, multiple sclerosis, Guillain-Barre syndrome, Crohn's disease, ulcerative colitis,

psoriasis, graft versus host disease, systemic lupus erythematosus, insulin-dependent diabetes mellitus, rheumatoid arthritis, toxic shock syndrome, Alzheimer's disease, diabetes, inflammatory bowel diseases, acute and chronic pain, stroke, myocardial infarction alone or following thrombolytic therapy, thermal injury, adult respiratory distress syndrome (ARDS), multiple organ injury secondary to trauma, acute glomerulonephritis, dermatoses with acute inflammatory components, acute purulent meningitis, syndromes associated with hemodialysis, leukopherisis, granulocyte transfusion associated syndromes, necrotizing enterocolitis, restenosis following percutaneous transluminal coronary angioplasty, traumatic arthritis, sepsis, chronic obstructive pulmonary disease and congestive heart failure, said method comprising administering to a patient a pharmaceutically effective amount of a compound according to claim 1.

17. A method of treating a disease or condition requiring anticoagulant or fibrinolytic therapy, said method comprising administering to a patient a pharmaceutically effective amount of a compound according to claim 1.

18. A process of making a compound of the formula (I):



20 Ar<sub>1</sub>, X, Y, Q, W, R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup> and R<sup>7</sup> are defined in claim 1;  
said process comprising

coupling under suitable conditions an amine bearing Ar<sup>1</sup> carboxylic acid of the formula (III), where P is a protecting group,

25

removing the protecting group P to provide an intermediate of formula (V) under suitable conditions;

coupling under suitable conditions the intermediate (V) with a halo heterocycle VI (Z = halogen) bearing R<sup>6</sup> in the presence of a suitable base to provide a compound of the formula (I):

